

## California Medical Device Recall Information



## **Recall Name**

## Bunnell Recalls Life Pulse High-Frequency Ventilator Patient Circuits Due to Possible Malfunction

Recall Date	Product Description	Recalling Firm	Recall Reason
12/12/12	Life Pulse High-Frequency Ventilator Patient Circuits	Bunnell, Inc. Salt Lake City, UT	Found to have heater wire insulation that can melt, causing sparking and smoke
Recall Class	Product Identification	Distribution	Affected Dates
I	List of Suspect Lots Recalled  (Note: Do not reference the Lot # on the outside of the circuit box. The recalled patient circuit lot # can be found on a label attached to the humidifier cartridge of each individual circuit.)	CA, nationwide and Canada	Distributed since March 19, 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm330842.htm